

Food Contact Declaration

FDA (Food & Drug Administration)

Product: PF CP 205 (Hp 2062.9)

We hereby declare that, according to the information provided by our raw material suppliers, the resin component (phenolic resin) of the product named below complies with the requirements of FDA regulation 21 CFR, Part 177, Section 2410 "Phenolic resins in molded articles".

Restrictions

- **No suitability for food contact:** PF CP 205 (Hp 2062.9) is a technical electrical-insulation phenolic paper laminate (paper-reinforced, phenolic-resin-bonded, flame-retardant special grade to UL 94 V-0) and, as a finished laminate, is NOT declared suitable for direct food contact.
- **Other:** Testing for extraction limits under FDA regulations has not been carried out on the finished laminate, since the product is intended for technical insulation applications (electrical, switchgear, mechanical engineering).

The exact provisions of the above-mentioned FDA regulation can be found at:
<https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/cfrsearch.cfm>

The quality management system of our raw material supplier is certified to DIN EN ISO 9001:2015 and forms an important basis for the consistent composition and quality of PF CP 205 (Hp 2062.9).

General note:

It remains the customer's responsibility to verify the suitability of plastic articles made from or with our products for their intended use in the food sector, i.e. for example:

- Verifying that the physical properties of the plastic are suitable for the intended applications,
- Verifying compliance with migration or extraction limits on the finished plastic parts,
- Verifying the possible influence of the plastic on the composition and/or organoleptic properties of foodstuffs.

Our statements are based on documents provided by our suppliers. No liability is assumed for the completeness or accuracy of the information contained herein. Existing laws and regulations must be observed by the recipient/user of our products on their own responsibility.

All information contained in this document is provided to the best of our knowledge and is based on sources considered reliable at the time of publication of this document. In the event of changes, for example due to legislation, manufacturing-related changes or new scientific findings, this declaration will be reassessed and becomes invalid. This declaration automatically expires 12 months after the date of issue. It is the sole responsibility of our customer to ensure compliance with the laws and regulations necessary for their intended use. Please request a current confirmation from Liedtke Kunststofftechnik if required.

The Food and Drug Administration (FDA) is the regulatory food safety and drug approval agency of the United States and reports to the Department of Health and Human Services.

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